FORM PTO-1449	U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE INFORMATION DISCLOSURE STATEMENT BY APPLICANT	ATTY. DOCKET NO. NNFF-1 CON	APPLN. NO. Not yet assigned
		APPLICANT Lan Kluwe	CONFIRMATION NO. Not yet assigned
		FILING DATE Concurrently Herewith	GROUP Not yet assigned

U.S. PATENT DOCUMENTS

EXAMINER INITIAL	DOCUMENT NUMBER	DATE	NAME	CLASS	SUBCLASS	FILING DATE IF APPROPRIATE
YK	6,077,685	06/20/00	Trofatter et al.	~~435~~	~~60.1~~	
	5,952,170	09/14/99	Stroun et al.	~~435~~	~~6~~	
	5,605,799	02/25/97	White et al.	~~435~~	~~6~~	
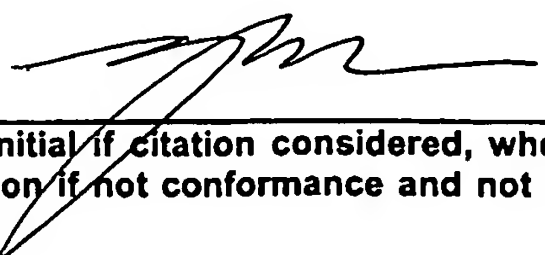
FOREIGN PATENT DOCUMENTS

EXAMINER INITIAL	DOCUMENT NUMBER	DATE	COUNTRY	CLASS	SUBCLASS	TRANSLATION	
						YES	NO

OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Pages, Etc.)

EXAMINER INITIAL	
YK	M.E. Baser, et al., "Presymptomatic diagnosis of neurofibromatosis 2 using linked genetic markers, neuroimaging, and ocular examinations," Neurology, 47:1269-1277 (1996).
	Irving et al., "Molecular Genetic Analysis of the Mechanism of Tumorigenesis in Acoustic Neuroma.," Arch. Otolaryngol. Head Neck Surg., 119:1222-1228 (1993).
	L. Kluwe, et al., "Mosaicism" in sporadic neurogibromatosis 2 patients," Human Molecular Genetics, 7(13):2051-2055 (1998).
	L. Kluwe, et al., "Allelic Loss of the NF1 Gene in NF1-Associated Plexiform Neurofibromas," Cancer Genet Cytogenet, 113:65-69 (1999).
	L. Kluwe, "Loss of NF1 Allele in Schwann Cells But Not in Fibroblasts Derived From an NF1-associated Neurofibroma," Genes, Chromosomes & Cancer, 24:283-285 (1999).

EXAMINER _____ DATE CONSIDERED 8-9-06

EXAMINER: Initial if citation considered, whether or not citation is in conformance with MPEP 609; Draw line through citation if not conformance and not considered. Include copy of this form with next communication to applicant.

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OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Pages, Etc.)

EXAMINER INITIAL		
YK		L. Kluwe, et al., "The parental origin of new mutations in neurofibromatosis 2," Neurogenetics, 3:17-24 (2000).
		L. Kluwe, et al., "Presymptomatic diagnosis for children of sporadic neurofibromatosis 2 patients: A method based on tumor analysis," Genetics in Medicine, 4(1):1-4 (2001).
		D.R. Lohmann, et al., "Molecular analysis and predictive testing in retinoblastoma," Ophthalmic Genetics, 16(4):135-142 (1995).
		V-F Mautner, et al., " Neurofibromatosis versus schwannomatosis", Fortschritte der Neurologie Psychiatrie, 66:271-277 (1998). [Abstract only - from BIOSIS Online, Biosciences Information Services, Philadelphia, PA].
		P. Riva, et al., "Characterization of a cytogenic 17q11.2 deletion in an NF1 patient with a contiguous gene syndrome," Hum Genet, 98:646-650 (1996).
		M. Sainio, et al., "Presymptomatic DNA and MRI diagnosis of neurofibromatosis 2 with mild clinical course in an extended pedigree," Neurology, 45:1314-1322 (1995).
		J. Sainz, et al., "Loss of Alleles in Vestibular Schwanomas," Archives of Otolaryngology-Head & Neck Surgery, 119:1285-1288 (1993).
		The 9th European Neurofibromatosis Meeting Program, April 6-8, 2001,Venice, Italy.
		K. Ueki et al., "Tight Association of Loss of Merlin Expression with Loss Heterozygosity at Chromosome 22q in Sporadic Meningiomas," Cancer Res., 59:5995-5998 (1999).
↓		Valero et al., "Linkage Disequilibrium Between Four Intragenic Polymorphic Microsatellites of the NF1 Gene and its Implications for Genetic Counselling," J. Mol. Genet., 3:590-593 (1996).

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